

1 Stephanie R. Tatar – State Bar No. 237792

2 **TATAR LAW FIRM, APC**

3 3500 West Olive Avenue

4 Suite 300

5 Burbank, California 91505

6 Tel. (323) 744-1146

7 Fax. (888) 778-5695

8 Stephanie@thetatarlawfirm.com

9

10 *Attorneys for Plaintiff*

11 *and the proposed Classes*

12

13 *Additional attorneys on signature page*

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16 **UNITED STATES DISTRICT COURT**
17
18 **NORTHERN DISTRICT OF CALIFORNIA**

19 SHOSHA KELLMAN, on behalf of herself and
20 all others similarly situated,

21 Plaintiff,

22 v.

23 WFM PRIVATE LABEL, L.P., WHOLE
24 FOODS MARKET CALIFORNIA, INC.,
25 WHOLE FOODS MARKET SERVICES, INC.
26 and WHOLE FOODS MARKET
27 DISTRIBUTION, INC.

Plaintiff,

Case No. 17-cv-06584-LB

**FOURTH AMENDED CLASS
ACTION COMPLAINT**

DEMAND FOR JURY TRIAL

Portion to Be Filed Under Seal
Pursuant to Order, Dkt. No. 89

Plaintiff Shosha Kellman, by her attorneys, brings this class action against WFM Private Label, L.P. (“WFM Private Label”), Whole Foods Market California, Inc. (“WFM California”), Whole Foods Market Services, Inc. (“WFM Services”), and Whole Foods Market Distribution,

1 Inc. (“WFM Distribution”) (collectively, “Defendants,” “WF” or “Whole Foods”), on her own
 2 behalf and on behalf of all others similarly situated, and alleges as follows:

3 **I. INTRODUCTION**

4 1. Whether an annoying patch of dry skin or an oozing rash that affects one’s social
 5 life, as much as 70% of the U.S. population is allergic to at least one personal care product
 6 ingredient. Most of these skin allergies are of unknown cause.

7 2. It is extremely difficult for people to identify what ingredient they are allergic to.
 8 Allergic reactions are attenuated in both space and time. Some allergic reactions will not manifest
 9 until a week after exposure to the allergen. Even worse – some allergic reactions will not manifest
 10 on the body part exposed to the allergen. Instead, the immune system will sometimes “remember”
 11 the first exposure and the allergic reaction will develop on the body part that was *first* exposed to
 12 the allergen.

13 3. Thus, consumers increasingly seek hypoallergenic products. Those who do not
 14 suffer from skin allergies seek hypoallergenic products to avoid developing a skin allergy. Those
 15 who do suffer from a skin allergy seek hypoallergenic products to avoid the inflammatory cascade
 16 caused by an unidentified skin allergen.

17 4. Since its founding, WF bases its brand as being a credible and trustworthy retailer,
 18 offering information and advice to consumers desiring safe products or seeking to avoid certain
 19 food ingredients or allergens.

20 5. In an effort to lure more customers, WF expanded to become not only a retailer and
 21 educator, but also a manufacturer of household and body care products. These private labels
 22 include 365 Everyday Value and Whole Foods Market product lines.

23 6. Seeking to capture the growing hypoallergenic market, WF prominently labels
 24 many of its products as “hypoallergenic.” *See* Product Labels attached as Exhibit 1.

25 7. However, despite its marketing scheme, WF’s products are chock-full of known
 26 skin sensitizers (allergens), agents that cause serious skin damage, chemicals that cause serious
 27 eye damage lasting longer than 21 days, skin irritants, and eye irritants. Even more, WF’s products

1 also contain known carcinogens, mutagens, reproductive toxins, and other chemicals extremely
2 hazardous to human health. *See Exhibit 1 and infra* at ¶¶ 106-134.

3 8. This is a class action on behalf of a national class of consumers who purchased
4 WF's body care products that were falsely and misleadingly marketed as "hypoallergenic." These
5 products in fact contain a shocking array of compounds known to cause allergic responses. These
6 products also contain a plethora of other compounds known to cause severe skin corrosion, serious
7 eye damage, or are otherwise toxic or hazardous in the case of skin contact. These products are
8 also stuffed with other chemicals that have not been analyzed for their skin sensitization potential.
9 Finally, these products also contain ingredients known to cause cancer, genetic mutations, birth
10 defects, or are otherwise toxic or hazardous to human health or the environment.

11 9. Many of the ingredients are permitted body care products. Yet WF did not simply
12 claim that its household products are "legal." WF falsely and misleadingly claimed that the
13 ingredients in its products are "hypoallergenic" when they are not.

14 10. By deceiving consumers about the nature, quality, and/or ingredients of its
15 products, WF is able to command a premium price, increasing consumers' willingness to pay and
16 take away market share from competing products, thereby increasing its own sales and profits.

17 11. Consumers lack the ability to test or independently ascertain the toxicity of a
18 chemical, especially at the point of sale. Reasonable consumers must and do rely on the chemicals
19 company to honestly report the nature of the product's ingredients.

20 12. WF further encouraged consumers to rely on its representations, marketing itself as
21 an honest company that provides transparent and truthful information about its products'
22 ingredients.

23 13. WF intended for consumers to rely on its representations, and hundreds of
24 thousands of reasonable consumers did in fact so rely.

25 14. As a result of its false and misleading labeling, WF was able to sell these products
26 to hundreds of thousands of consumers throughout the United States and to profit handsomely
27 from these transactions.

1 15. WF's false and misleading representations and omissions violate state and federal
2 law, both civil and criminal, detailed more fully below, including California's Unfair Competition
3 Law, California's Consumer Legal Remedies Act, and common law.

4 16. Plaintiff brings this action to stop WF's deceptive and misleading practices.

II. PARTIES

A. Plaintiff Shosha Kellman

7 17. Plaintiff Shosha Kellman is an individual consumer who, at all times material
8 hereto, was a citizen of the State of California and resident of Alameda. For approximately twenty-
9 four months, from early 2014 through early 2016, Plaintiff Kellman regularly purchased WF's 365
10 Gentle Skin Cleanser from the Whole Foods Market located at 3000 Telegraph Ave, Berkeley CA
11 94705 and from the Whole Foods Market located at 230 Bay Place, Oakland, CA 94612. Ms.
12 Kellman consistently used a credit card for her purchases. Plaintiff Kellman estimates that she
13 purchased the product every four to six weeks. In addition, Plaintiff Kellman purchased other WF
14 products. Plaintiff Kellman sometimes purchased WF's 365 moisturizing lotion during the same
15 24-month period of time.

16 18. In deciding to make these purchases, Plaintiff Kellman saw, relied upon, and
17 reasonably believed the label representation that the products were "hypoallergenic." These
18 representations were a significant reason for her purchases.

19 19. Plaintiff Kellman and her family members have all suffered skin irritation, eye
20 irritation, dermatitis, and/or an allergic skin reaction in the past.

21 20. In the case of common skin irritation or dermatitis, Plaintiff Kellman, like similarly
22 situated consumers, is unsure whether what seemed like skin or eye irritation or dermatitis was in
23 fact an allergic response to an ingredient in a personal care product.

24 21. Like similarly situated consumers, Plaintiff Kellman does not know the identity of
25 every ingredient she and her family are allergic to. Moreover, like similarly situated consumers,
26 Plaintiff Kellman does not know which ingredients she or her family may develop an allergy to.

27 22. Had Plaintiff Kellman known at the time that these products were not

1 hypoallergenic as promised, she would not have purchased these products.

2 23. Had Plaintiff Kellman known at the time that these products contained irritating,
3 toxic, hazardous, or otherwise harmful chemicals, she would not have purchased these products.

4 24. Plaintiff Kellman purchased, purchased more of, or paid more for, these products
5 than she would have had she known that the products contained skin sensitizers, irritants, toxins,
6 carcinogens, or otherwise harmful chemicals.

7 25. If WF's products were reformulated such that its representations were truthful,
8 Plaintiff Kellman would consider purchasing WF's products in the future.

9 26. The products that Plaintiff Kellman purchased are substantially similar to WF's
10 other products alleged to be falsely labeled.

11 **B. Defendant WFM Private Label, L.P.**

12 27. Defendant WFM Private Label, L.P. ("WFM Private Label") is a Delaware
13 corporation, doing business in the State of California and throughout the United States of America.
14 See, e.g., Exhibit 2.

15 28. WFM Private Label is a wholly owned subsidiary of Whole Foods Market, Inc. and
16 an affiliate of Whole Foods Market California, Inc., Whole Foods Market Services, Inc., and
17 Whole Foods Market Distribution, Inc.

18 **C. Defendant Whole Foods Market California, Inc.**

19 29. Whole Foods Market California, Inc. ("WFM California") is a California
20 corporation and operates the Whole Foods retail stores in Northern California.

21 **D. Defendant Whole Foods Market Services, Inc.**

22 30. Whole Foods Market Services, Inc. ("WFM Services") controls decisions relating
23 to the nation-wide design, development, advertising, and marketing of Whole Foods private label
24 lines of products. *See Exhibit 3, Decl. of WFM Services Private Label – Customer Service and*
25 *Sales Support Team Leader, Rebecca Stuch, Frame v. Whole Foods Market, Inc., Case No. 1:06-*
26 *cv-07058-DAB ("Frame")*, ECF 6 at ¶ 4 (WFM Services Private Label – Customer Service and
27 Sales Support Team Leader declaring that "All national marketing and advertising for the 365

1 EVERYDAY VALUE and 365 ORGANIC EVERYDAY VALUE line of products is directed
2 from Whole Foods Market Services, Inc.’s offices in Austin, Texas.”); Exhibit 4, Excerpts from
3 Memorandum of Law in Support of Motion to Transfer, *Frame* ECF 8 at 2 (“Decisions relating to
4 the design, development, advertising, and marketing of these private label lines of products are
5 controlled by employees of Whole Foods Market Services, Inc.”).

6 **E. Defendant Whole Foods Market Distribution, Inc.**

7 31. Whole Foods Market Distribution, Inc. (“WFM Distribution”) is a Delaware
8 corporation that distributes or causes the distribution of all WF’s private label products throughout
9 the United States, including and specifically to retail stores in the State of California. *See* Exhibit
10 5, Excerpts from Agreement for the Distribution of Products between WFM Distribution and
11 United Natural Foods, Inc.

12 **III. JURISDICTION AND VENUE**

13 32. This Court has personal jurisdiction over the parties in this case.

14 **A. Plaintiff**

15 33. Plaintiff Kellman is a citizen of California.

16 **B. Defendants**

17 34. This Court has personal jurisdiction over Defendants because they have sufficient
18 minimum contacts in California or otherwise intentionally avail themselves of the laws of this
19 State through the marketing of the products at issue in California to consumers in California and
20 through direct sales of the products at issue in California to consumers in California, so as to render
21 the exercise of jurisdiction by this Court consistent with traditional notions of fair play and
22 substantial justice.

23 **C. Additional Allegations Of Personal Jurisdiction Over Defendants**

24 35. WFM Services, WFM California, WFM Distribution, and WFM Private Label all
25 use common marketing, including a common use of Whole Foods trademarks and logos. WF uses
26 this common, nation-wide marketing scheme to give consumers the belief that its stores have a
27

1 national presence and abide by a nation-wide set of standards.

2 36. WF uses this nation-wide marketing scheme to give consumers the belief that all
3 Whole Foods entities are, in practicality, one organization. WF uses a nation-wide marketing
4 scheme to convey to consumers that the Whole Foods retailer that sells natural and organic
5 products is also producing the private label Whole Foods personal care and household cleaning
6 products.

7 37. Thus, Whole Foods retailers are the exclusive retailer for Whole Foods personal
8 care and household cleaning products.

9 38. WFM California is incorporated in California. WFM California operates the Whole
10 Foods retail stores in certain California regions, including northern California.

11 39. WFM Distribution is the exclusive distributor of WF's private label products in
12 certain California regions. *See Exhibit 5.*

13 40. WFM Private Label is the Whole Foods entity responsible for Whole Foods private
14 label products and exclusive brands.

15 41. Whole Foods private label products are sold in Whole Foods Market retail locations
16 in the United States, some of which are owned and operated by WFM California.

17 42. WFM Private Label contracts with vendors to produce Whole Foods private label
18 products, which are then sold throughout the United States, including to Whole Foods retail
19 locations in California. Specifically, WFM Private Label enters into contracts with vendors
20 whereby vendors agree to supply Whole Foods private label products to stores and distribution
21 centers owned or operated by WFM Private Label or its affiliates. An example of such a vendor
22 agreement is attached hereto as Exhibit 2.

23 43. WFM Private Label has a vendor agreement that is substantially similar to Exhibit
24 2 for all products that are included in the putative class, and specifically for all products that are
25 sold in California.

26 44. Pursuant to its vendor contracts, WFM Private Label retains full approval rights
27 and final authorization with regard to private label product packaging, labeling, and formulation,

1 and vendors cannot order or produce labels or packaging for private label products until WFM
2 Private Label has provided written approval and authorization. Exhibit 2.

3 45. WFM Private Label regularly performs multiple acts purposefully directed to
4 California, for the purpose and with the intent that WF private label products continue to be sold
5 in California.

6 46. For example, WFM Private Label specifically negotiated for vendors to indemnify
7 it for any claim relating to the California Safe Drinking Water and Toxic Enforcement Act of 1986
8 (Proposition 65), thereby specifically contemplating for the agreement to be performed at least
9 partially in California. Exhibit 2 at § 11(a) at WFM00097.

10 47. As another example, the California Air Resources Board regulates consumer
11 products that are a significant source of volatile organic compounds and requires each company
12 listed on the label of a consumer product sold in California to complete the annual Consumer &
13 Commercial Products Survey.

14 48. To maintain the ability to sell WF private label products, WFM Private Label
15 completed the annual Consumer & Commercial Products Survey, submitting to the California Air
16 Resources Board information regarding its laundry detergents, body washes, lotions, and shampoo
17 (i.e., all products included in this suit). *See, e.g.*, Draft List of Companies and Number of Products
18 Reported in 2014, *available for download at* <https://preview.tinyurl.com/WFMPL-CARB>.

19 49. WFM Private Label forbids vendors from selling any Whole Foods private label
20 products to anyone without WFM Private Label's approval. *See, e.g.*, Exhibit 2 at § 10.

21 50. Thus, WFM Private Label authorized the sale of all WF private label products sold
22 in California.

23 51. WFM Private Label also retains the right to alter or change any private label product
24 packaging or labeling that does not conform to contract specifications or requirements that WFM
25 Private Label provides. Exhibit 2.

26 52. The labels for which WFM Private Label retains final authority and approval
27 include products obtained from vendors that supply WFM Private Label affiliates that operate and

1 sell such products in California, including WFM California and the subject products outlined
 2 below.

3 53. WFM Services is the marketing arm of Whole Foods, and Whole Foods subsidiaries
 4 are the exclusive retailers of WFM private label products.

5 54. WFM Services designs, develops, advertises, and markets Whole Foods private
 6 label products to be sold in California.

7 55. WFM Services accomplishes the national design, development, advertising, and
 8 marketing of its private label lines of products through, *inter alia*, its assets and employees based
 9 in the State of California.

10 56. In 2011, WFM Services acquired all assets of WFM Brand 365 LLC, the former
 11 “private label” entity for WF stores. Exhibit 6 (transferring all assets of WFM Brand 365 LLC to
 12 WFM Services).

13 57. WFM Services owns and operates Whole Foods’ website, which is used to market
 14 and consummate online sales of its products to consumers in California and throughout the nation.
 15 *See, e.g.*, Whole Foods Privacy Notice, <https://www.wholefoodsmarket.com/privacy-policy> (last
 16 visited Feb. 15, 2018), and Whole Foods Terms & Conditions,
 17 <https://www.wholefoodsmarket.com/terms-use> (last visited Feb. 15, 2018).

18 58. WFM Services sells Whole Foods private label products at issue in this suit to
 19 California customers through the Whole Foods website, which WFM Services owns and operates.

20 59. Through the Whole Foods website, WFM Services directs customers to purchase
 21 WF private label products through the Whole-Foods specific page at the California-based Instacart.

22 60. WFM Services has enjoyed substantial revenues through its online sales to
 23 California consumers. More specifically, WFM Services has enjoyed substantial revenues through
 24 its online sales to California consumers of WF private label products at issue in this suit.

25 61. Online, WFM Services advertises “local” deals specific to California customers,
 26 and its online “store locator” also helps California consumers locate the California-based retailer
 27 nearest them. It also provides notices specifically to California residents to further facilitate

1 commercial activities in California, such as notices regarding credit card investigations related to
2 payments they made at California locations, and notices of their privacy rights under California
3 law.

4 62. WFM Services has continuous and systematic general business contacts that
5 approximate physical presence in California and many of its activities are specifically directed to
6 California.

7 63. WFM Services works in creating the design of WF's California stores and planning
8 their dates of opening. WFM Services also assists WF's California stores with their point-of-sale
9 systems and with auditing the vendors who manufacture products for WF. In addition to
10 performing accounting, legal, and other administrative services for WF's California stores, WFM
11 Services also develops policies and the General Information Guidelines¹ with the various WF
12 regions.

13 64. WFM Services also plays a key role in the WF private label products at issue in this
14 suit. It is responsible for marketing Whole Foods brand. WFM Services is also responsible for
15 marketing WF's private label products online.

16 65. The "Quality Standards Team" at WFM Services determines which ingredients,
17 vendors, and product types are available to WF's California retailers.

18 66. [REDACTED]

19 [REDACTED]

20 [REDACTED]

21 _____

22 ¹ The General Information Guidelines (a.k.a. "GIG") states the terms and conditions of WF
23 subsidiaries' relationships with its employees, including pay, work conditions, training, work
24 requirements, non-competition provisions, confidentiality requirements, and various employment
25 policies such as dress code, recordings in the workplace, smoking policies, and policies for
26 communicating in online forums such as internet chat rooms and message boards. *See* WF Code
27 of Business Conduct 2017, [http://s21.q4cdn.com/118642233/files/doc_downloads/governance_documents/2017/Whole-Foods-Market-Code-of-Business-Conduct\[1\].pdf](http://s21.q4cdn.com/118642233/files/doc_downloads/governance_documents/2017/Whole-Foods-Market-Code-of-Business-Conduct[1].pdf); WF and Local 919, recordings in workplace, <http://hr.cch.com/ELD/WholeFoods122415.pdf>; WF Executive Retention Plan, <https://www.sec.gov/Archives/edgar/data/865436/000086543612000017/wfmexecutiveretentionplane.htm>

1 [REDACTED]
2 [REDACTED]
3 **D. Subject Matter Jurisdiction**

4 67. This Court has subject matter jurisdiction pursuant to the Class Action Fairness
5 Act, 28 U.S.C. § 1332(d) (“CAFA”). Jurisdiction under CAFA is met because the proposed
6 number of putative class members exceeds 100, at least one plaintiff and one defendant are citizens
7 of different states, and the amount in controversy, including, but not limited to the aggregate
8 amount of relief sought by absent class members, exclusive of interest and costs, exceeds \$5
9 million.

10 68. Venue is proper in this District under 28 U.S.C. § 1331(a). Substantial acts in
11 furtherance of the alleged improper conduct, including the dissemination of false, misleading and
12 deceptive information regarding the nature, quality, and/or ingredients of the products, occurred
13 within this District.

14 69. No other forum would be more convenient for the parties and witnesses to litigate
15 this action.

16 **III. FACTUAL ALLEGATIONS**

17 **A. Consumers Actively Seek Hypoallergenic Body Care Products**

18 70. According to the Centers for Disease Control and Prevention (“CDC”), 8.8 million
19 children (12% of U.S. children) reported skin allergies in 2012. Skin allergies are even more
20 prevalent among young children; CDC reports that 14.2% of children between the ages of 0 and 4
21 suffered a skin allergy in 2012.

22 71. These numbers are likely to underreport the prevalence of allergic contact
23 dermatitis; recent studies show that somewhere between 14-70% of children suffer from skin
24 allergies, based on positive patch skin tests.

25 72. Skin allergies are similarly prevalent among adults.

26 73. When skin is exposed to a sufficient amount of a chemical allergen, the skin is
27 “sensitized.” Upon re-exposure to the allergen, the skin initiates an inflammatory cascade, causing

1 skin changes associated with allergic contact dermatitis. These include redness, oedema (fluid
2 retention), scaling, fissures (cracking), vesicles (fluid-filled sacs), bullae (bubble-like cavity), and
3 eventually oozing.

4 74. Contact sensitization and related skin allergies can severely affect a person's quality
5 of life, depending on the severity and the site of skin sensitization. People suffering from
6 noticeable skin allergies will try to hide the symptoms under clothing if possible, and if not, will
7 avoid public spaces entirely. In either case, skin allergies can dramatically affect a person's
8 confidence and engagement in life.

9 75. It is difficult to identify the substance causing an allergic response. Allergic contact
10 dermatitis develops several days after exposure to a skin allergen. Some substances do not cause
11 symptoms until a week after exposure.

12 76. Even more, once an individual is sensitized to an allergen, future contact with the
13 allergen can trigger a response in the *original* site of sensitization. For example, if someone had
14 an allergic response to a product used on the face, and later used a different product containing the
15 same allergen on the legs, the allergic response will occur again on the *face* – even if the face was
16 never exposed to the second product.

17 77. When a consumer cannot identify the material to which they are allergic, allergic
18 contact dermatitis will persist, and, it is believed, will take longer to resolve even after the cause
19 is identified.

20 78. Thus, consumers will actively seek out hypoallergenic products – to avoid a skin
21 allergy from occurring at all and/or to prevent a known skin allergy from repeating the
22 inflammatory cascade.

23 **B. Definition of Hypoallergenic**

24 79. The scientific and regulatory definition of a skin sensitizer is a substance that causes
25 sensitization by skin contact in a substantial number of persons based on human evidence or
26 appropriate animal testing.

27 80. If a skin sensitizer makes up 0.1% or more of a product, or if the product contains

1 a sensitizer that may elicit an allergic response at concentrations smaller than 0.1% in individuals
2 who are already sensitized to the chemical, the *entire* product mixture is classified as a skin
3 sensitizer, *i.e.*, the product causes sensitization by skin contact in a substantial number of persons
4 based on human evidence or appropriate animal testing.

5 81. A product that is a skin sensitizer is not hypoallergenic.

6 82. Consumers believe and expect that a product that is labeled as hypoallergenic does
7 not contain skin sensitizers at a concentration that could elicit an allergic response in sensitized
8 individuals.

9 83. Once skin is sensitized, even a *minute* amount of the chemical allergen is enough
10 to cause a full-blown allergic response. Thus, consumers seeking hypoallergenic products also
11 commonly expect that the product does not contain *any* skin sensitizers.

12 84. All WF's products contain substances classified by reputable authorities as skin
13 sensitizers. *See infra* at ¶¶ 106-134 (identifying skin sensitizers) and Exhibit 1 (showing which
14 products contain these skin sensitizers).

15 85. All WF's products also contain skin sensitizers that are either present in WF's
16 products at concentrations larger than 0.1%, or that may elicit an allergic response at
17 concentrations smaller than 0.1% in sensitized individuals.

18 86. Thus, WF's products are not hypoallergenic.

19 87. Thus, WF's on-the-label promise that its products are "hypoallergenic" is false.

20 88. Consumers also believe and expect that a hypoallergenic product will not cause
21 skin irritation, skin corrosion, or eye damage when used as directed.

22 89. Consumers also believe and expect that a product that is labeled as hypoallergenic
23 does not contain a significant amount of ingredients known to produce skin irritation, skin
24 corrosion, and/or eye damage.

25 90. WF's products contain significant amounts of ingredients classified by reputable
26 authorities as causing skin irritation, skin corrosion, and/or eye damage. *See infra* at ¶¶ 106-134
27 and Exhibit 1 (showing which products contain these ingredients).

1 91. Thus, WF's on-the-label promise that its products are "hypoallergenic" is *also*
2 misleading.

3 92. WF knows how consumers understand "hypoallergenic," and encourages this
4 understanding.

5 93. Because even a *minute* amount of a chemical allergen is enough to cause a full-
6 blown allergic response, consumers reasonably expect and believe that when a product is labeled
7 as "hypoallergenic," this representation is true not just for the final formulation, but to every
8 ingredient in the product.

9 94. WF knows and encourages this understanding.

10 95. WF knows that consumers rely upon it to not only test the final product formulation
11 for basic safety, but to select only those ingredients that it considers to be safe.

12 96. Advertising itself as "America's Healthiest Grocery Store," *see* Exhibit 9 (Google
13 ad); 2016 Annual Report at 1, Whole Foods promises its customers that it "maintain[s] the strictest
14 quality standards in the industry." Exhibit 10 ("Company Info").

15 97. Listing its "quality standards," Whole Foods identifies as its top standard: "We
16 carefully evaluate each and every product we sell." Exhibit 11 ("Quality Standards").

17 98. WF stresses not only product safety, but *ingredient* safety. As WF explains:

18 **OUR BODY CARE QUALITY STANDARDS**

19 We carry the finest, high-quality beauty, hair and body care products
20 available because we believe the quality of the items and ingredients you put on
your body is as important as the foods and nutritional supplements you put in your
21 body. We evaluate the quality of personal care products in terms of ingredients,
experience, and efficacy.

22 Exhibit 12 ("Body Care Quality Standards").

23 99. However, many ingredients in WF's products have not been adequately studied for
24 safety. Moreover, very few have been assessed for their sensitization potential. *See ¶¶ 106-134,*
25 *infra.*

1 C. **WF's False Representations**
2

3 100. On the products' labels, and again on its retail website, WF represents that certain
4 of its products are "hypoallergenic." These products, (collectively, the "False Labeled Products")
5 are all falsely labeled, as all of these products contain skin sensitizers, skin irritants, eye irritants,
6 and other deleterious compounds.
7

8 101. These products are:
9

10 365 Baby Foaming Wash
11 365 Baby Lotion
12 365 Baby Shampoo
13 365 Bubble Bath
14 365 Gentle Skin Cleanser
15 365 Kids' Foaming Wash
16 365 Maximum Moisture Body Lotion
17 365 Moisturizing Lotion
18 Whole Foods Market Baby Laundry Detergent
19 Whole Foods Market Organic Laundry Detergent
20 Wild Kratts Bubble Bath
21 Wild Kratts Kids Foaming Body Wash

22 102. The labels of these products are attached as Exhibit 1.

23 103. Further encouraging consumers' reliance on WF's "hypoallergenic" promise, WF
24 labels only *some* products as hypoallergenic, giving consumers the (false) impression that WF
25 carefully reviewed each ingredient in its products to ensure that the "hypoallergenic" promise was
26 made for only those products that truly are hypoallergenic. *See, e.g.*, Exhibit 13.

27 104. Yet, contrary to WF's promise, *all* these products in fact contain known skin
28 sensitizers. They also *all* contain known skin or eye irritants, carcinogens, teratogens, mutagens,
29 or pollutants. Finally, they *all* contain substances that have not been adequately assessed for safety
30 or skin sensitization potential.

31 105. All WF's False Labeled Products contain one or more of the following chemicals.

32 106. *Acacia senegal (organic gum arabic)* is classified as a Category 1 skin sensitizer,
33 based on positive animal and/or human testing demonstrating that repeated skin contact can be
34 expected to cause an allergic response in a substantial number of persons. It is known to cause
35 local contact dermatitis. It is a Category 2 skin irritant, meaning that it causes significant
36 erythema/eschar (redness and dead tissue) or edema (abnormal accumulation of fluid beneath the

1 skin) lasting more than three days, or skin inflammation lasting longer than 14 days. It is a
 2 Category 2 eye irritant, causing adverse effects on the cornea, iris, and conjunctiva.

3 107. Some testing classifies *calendula officinalis flower extract* as a Category 1 skin
 4 sensitizer, based on positive animal and/or human testing demonstrating that repeated skin contact
 5 can be expected to cause an allergic response in a substantial number of persons. It is a Category
 6 2 skin irritant, meaning that it causes significant erythema/eschar (redness and dead tissue) or
 7 edema (abnormal accumulation of fluid beneath the skin) lasting more than three days, or skin
 8 inflammation lasting longer than 14 days. It is a Category 2 eye irritant, causing adverse effects
 9 on the cornea, iris, and conjunctiva.

10 108. *Caprylyl glycol* causes Category 1 eye damage, i.e., it causes serious damage to the
 11 eye tissue or serious physical decay of vision which is not fully reversible within 21 days of
 12 application.

13 109. *Cetearyl alcohol* is recognized as an allergen by the American Contact Dermatitis
 14 Society. Its safety for use in bodycare products has not been adequately assessed. The limited
 15 testing done, however, shows it to be a skin irritant and eye irritant, causing skin damage in less
 16 than four hours and adverse effects on the cornea, iris, conjunctiva. It is inherently toxic to aquatic
 17 life. It is also toxic to the mucous membranes and is hazardous by definition under federal law.

18 110. *Cetyl alcohol* has caused urticaria-like dermatitis in humans. It is also a skin and
 19 eye irritant. It is also classified as an eye irritant, and it is inherently toxic to aquatic life with long-
 20 lasting effects.

21 111. While *citric acid* is a common food ingredient, skin contact is known to cause
 22 allergic reactions in humans. It has been reported to cause Category 1B skin corrosion, meaning
 23 that it irreversibly damages the skin after short exposure; in animal tests, the substance caused
 24 visible necrosis after less than 1 hour of exposure. Corrosive reactions are typified by ulcers,
 25 bleeding, bloody scabs, and, by the end of observation at 14 days, by discoloration due to blanching
 26 of the skin, complete areas of alopecia, and scars. It causes Category 1 eye damage, i.e., it causes
 27 serious damage to the eye tissue or serious physical decay of vision which is not fully reversible

1 within 21 days of application.

2 112. Repeated use of *cocamidopropyl hydroxysultaine* has caused increased skin
 3 irritation. In one test on human subjects, while no skin irritation was observed at the first
 4 application of a 2.5% solution of cocamidopropyl hydroxysultaine, repeated applications caused
 5 slight to moderate skin irritation in 45% of the subjects, with 5% of the subjects developing strong
 6 irritation. It causes Category 1 eye damage, *i.e.*, it causes serious damage to the eye tissue or
 7 serious physical decay of vision which is not fully reversible within 21 days of application.

8 113. *Decyl glucoside* has caused sensitization in human testing and is recognized as an
 9 allergen by the American Contact Dermatitis Society. It causes Category 1C skin corrosion,
 10 meaning that it irreversibly damages the skin after short exposure; in animal tests, the substance
 11 caused visible necrosis after less than 4 hours of exposure. Corrosive reactions are typified by
 12 ulcers, bleeding, bloody scabs, and, by the end of observation at 14 days, by discoloration due to
 13 blanching of the skin, complete areas of alopecia, and scars. It causes Category 1 eye damage, *i.e.*,
 14 it causes serious damage to the eye tissue or serious physical decay of vision which is not fully
 15 reversible within 21 days of application.

16 114. The sensitization potential of *gluconolactone* has not been assessed by any
 17 reputable authority. However, based on its chemical structure and similarity to other known skin
 18 sensitizers, it is classified as a likely skin sensitizer.

19 115. *Glycerin (also listed as "organic glycerin")* is known to cause eczema in humans.
 20 Based on its chemical structure and similarity to other known skin sensitizers, it is a suspected skin
 21 sensitizer. Glycerin (also listed as "organic glycerin") is classified as a skin and eye irritant. It is a
 22 mutagen, meaning that it is suspected of mutating human cells in a way that can be transmitted to
 23 children conceived after exposure.

24 116. *Glyceryl stearate* is a skin and eye irritant. In animal testing (rabbits), it caused
 25 erythema, edema, atonia, desquamation, and/or fissuring. It is also inherently toxic to aquatic life.

26 117. *Isopropyl palmitate* is classified as a skin and eye irritant. Moreover, it is an ester,
 27 a class of chemicals known to be environmentally toxic.

1 118. Some testing classifies *mentha viridis (spearmint) leaf oil* as a Category 1 skin
 2 sensitizer, based on positive animal and/or human testing demonstrating that repeated skin contact
 3 can be expected to cause an allergic response in a substantial number of persons. *Mentha viridis*
 4 (*spearmint*) leaf oil is classified as a fragrance allergen in the European Union. It is a Category 2
 5 skin irritant, meaning that it causes significant erythema/eschar (redness and dead tissue) or edema
 6 (abnormal accumulation of fluid beneath the skin) lasting more than three days, or skin
 7 inflammation lasting longer than 14 days. It is a Category 2 eye irritant, causing adverse effects
 8 on the cornea, iris, and conjunctiva.

9 119. WF does not disclose the identity of the fragrances it uses, listing only the generic
 10 term “*natural fragrance*” on its product label. Many synthetic fragrances are known to be human
 11 sensitizers, toxins and environmental hazards, and are associated with adverse reproductive
 12 effects, genetic mutations, and other ill effects. As WF itself recognizes, “[p]hthalates have been
 13 linked to cancer and endocrine system disruption and are currently covered under the umbrella
 14 term “fragrance” in conventional products.” Exhibit 14 (“What You Won’t Find in Our Cleaning
 15 Products”).

16 120. *Olea europaea (olive) oil* is classified as a skin irritant. (Thus, masseurs are
 17 discouraged from the external use of olive oil). It is also classified as an eye irritant.

18 121. The sensitization potential of *panthenol* has not been assessed by any reputable
 19 authority. However, based on its chemical structure and similarity to other known skin sensitizers,
 20 it is a likely skin sensitizer. In fact, it has produced allergic responses in some past testing on
 21 humans. Panthenol is classified as a skin and eye irritant.

22 122. *Phenoxyethanol* is a skin and severe eye irritant. It has induced an allergic response
 23 in both human and animal testing. It is recognized as an allergen by the American Contact
 24 Dermatitis Society. It is toxic by all routes (inhalation, ingestion, and dermal contact). It is
 25 extremely hazardous in case of eye contact and very hazardous in case of skin contact (defatting
 26 the skin and causing skin inflammation characterized by itching, scaling, reddening, or,
 27 occasionally, blistering). Even short exposure can cause serious temporary or residual injury. It

1 is toxic to the kidneys, the nervous system, and the liver, adversely affecting the central nervous
 2 system and peripheral nervous system, causing headaches, tremors, and central nervous system
 3 depression. It degrades into substances that are even more toxic. It is a germ cell mutagen,
 4 suspected of mutating human cells in a way that can be transmitted to children conceived after
 5 exposure. It is also a reproductive toxin, suspected of damaging fertility or the unborn child based
 6 on human or animal evidence. Phenoxyethanol is an ethylene glycol ether, which is known to cause
 7 wasting of the testicles, reproductive changes, infertility, and changes to kidney function.
 8 Phenoxyethanol is also carcinogen, meaning that it is suspected to induce cancer or increase its
 9 incidence. Case studies indicate that repeated exposure to phenoxyethanol results in acute
 10 neurotoxic effects, as well as chronic solvent-induced brain syndrome, constant irritability,
 11 impaired memory, depression, alcohol intolerance, episodes of tachycardia and dyspnea, and
 12 problems with balance and rash. Phenoxyethanol is also toxic by definition under federal law, and
 13 is regulated as a toxic compound. Its use is restricted in Europe.

14 123. **Polysorbate 20** is classified as a Category 1 skin sensitizer, based on multiple
 15 positive tests demonstrating that repeated skin contact can be expected to cause allergic response
 16 in a substantial number of persons. It is also a Category 2 skin and eye irritant, causing skin damage
 17 in less than four hours and adverse effects on the cornea, iris, and conjunctiva. It is made in part
 18 with ethylene oxide, resulting in 1,4 dioxane as a trace contaminant, which is classified as a
 19 possible carcinogen. It is a teratogen, meaning that it causes birth defects.

20 124. **Polysorbate 60** has caused urticaria (hives and swelling) on human subjects'
 21 foreheads. In animal testing, polysorbate 60 is a skin irritant.

22 125. The sensitization potential of **potassium sorbate** has not been assessed by any
 23 reputable authority. However, based on its chemical structure and similarity to other known skin
 24 sensitizers, it is classified as a likely skin sensitizer. Some case studies show it to cause contact
 25 urticaria. It is a Category 2 skin irritant, meaning that it causes significant erythema/eschar
 26 (redness and dead tissue) or edema (abnormal accumulation of fluid beneath the skin) lasting more
 27 than three days, or skin inflammation lasting longer than 14 days. Some studies show it to cause

1 Category 1A skin corrosion, meaning that it irreversibly damages the skin after short exposure; in
2 animal tests, the substance caused visible necrosis after less than 3 minutes of exposure. Corrosive
3 reactions are typified by ulcers, bleeding, bloody scabs, and, by the end of observation at 14 days,
4 by discoloration due to blanching of the skin, complete areas of alopecia, and scars. It is a
5 Category 2 eye irritant, causing adverse effects on the cornea, iris, and conjunctiva. It is also a
6 suspected mutagen.

7 126. Some testing classifies **sodium benzoate** as a Category 1 skin sensitizer, based on
8 positive animal and/or human testing demonstrating that repeated skin contact can be expected to
9 cause an allergic response in a substantial number of persons. It is also a skin irritant and causes
10 serious eye damage. It is a Category 2 eye irritant, causing adverse effects on the cornea, iris, and
11 conjunctiva. Some testing finds that it causes Category 1 eye damage, i.e., it causes serious damage
12 to the eye tissue or serious physical decay of vision which is not fully reversible within 21 days of
13 application. It is a teratogen, meaning that it causes birth defects. Its use in personal care products
14 is limited in Europe.

15 127. **Sodium bicarbonate** is classified as a skin and eye irritant. Some tests show that it
16 causes Category 1 eye damage, i.e., it causes serious damage to the eye tissue or serious physical
17 decay of vision which is not fully reversible within 21 days of application. It is a teratogen,
18 meaning that it causes birth defects.

19 128. **Sodium carbonate** is a skin and eye irritant. It causes Category 1 eye damage, i.e.,
20 it causes serious damage to the eye tissue or serious physical decay of vision which is not fully
21 reversible within 21 days of application.

22 129. The sensitization potential of **sodium citrate** has not been assessed by any reputable
23 authority. However, based on its chemical structure and similarity to other known skin sensitizers,
24 it is classified as a suspected skin sensitizer. It is also classified as a skin and eye irritant, causing
25 significant erythema/eschar (redness and dead tissue) or edema (abnormal accumulation of fluid
26 beneath the skin) lasting more than three days, or skin inflammation lasting longer than 14 days,
27 and causing adverse effects on the cornea, iris, and conjunctiva.

1 130. The sensitization potential of *sodium myristoyl sarcosinate* has not been assessed
 2 by any reputable authority. However, based on its chemical structure and similarity to other known
 3 skin sensitizers, it is classified as a suspected skin sensitizer. It is a Category 2 skin irritant,
 4 meaning that it causes significant erythema/eschar (redness and dead tissue) or edema (abnormal
 5 accumulation of fluid beneath the skin) lasting more than three days, or skin inflammation lasting
 6 longer than 14 days. It causes Category 1 eye damage, i.e., it causes serious damage to the eye
 7 tissue or serious physical decay of vision which is not fully reversible within 21 days of
 8 application.

9 131. The sensitization potential of *sodium oleate* has not been assessed by any reputable
 10 authority. However, based on its chemical structure and similarity to other known skin sensitizers,
 11 it is classified as a suspected skin sensitizer.

12 132. Though *xanthan gum* is safe as a food ingredient, it is not so safe for the
 13 skin. Some testing indicates that it is a skin sensitizer. It is a Category 2 skin irritant, meaning
 14 that it causes significant erythema/eschar (redness and dead tissue) or edema (abnormal
 15 accumulation of fluid beneath the skin) lasting more than three days, or skin inflammation lasting
 16 longer than 14 days. It is a Category 2 eye irritant, causing adverse effects on the cornea, iris, and
 17 conjunctiva.

18 133. *Cyamopsis tetragonolobus gum (organic guar gum)* is a contact
 19 sensitizer. Additionally, it is a Category 2 eye irritant, causing adverse effects on the cornea, iris,
 20 and conjunctiva.

21 134. *Avena sativa (oat) kernel flour, or avena sativa kernel flour* is classified as a
 22 Category 1 skin sensitizer, based on positive animal and/or human testing demonstrating that
 23 repeated skin contact can be expected to cause an allergic response in a substantial number of
 24 persons.

25 **D. The Representations Are False, Deceptive, And Misleading**

26 135. WF's conduct deceived and/or was likely to deceive the public. Consumers were
 27 deceived into believing that the Falsey Labeled Products were hypoallergenic, as labeled.

136. All these representations were false, as explained *supra*.

137. Consumers would not know the true nature of the ingredients merely by reading the ingredient label. Its discovery requires investigation beyond the grocery store and knowledge of chemistry beyond that of the average reasonable consumer.

5 **E. Location Of The Misrepresentations**

6 138. WF made the above false, deceptive, and misleading misrepresentations and
7 omissions on the package of the Falsely Labeled Products. *See* Exhibit 1.

8 139. WF repeated the above false, deceptive, and misleading misrepresentations and
9 omissions on its online retail product page for the Falsely Labeled Products. *See* Exhibit 1.

10 140. The misrepresentations and omissions were uniform and have actually been
11 communicated to Plaintiff and to each member of the Class at every point of purchase and
12 consumption.

13 **F. WF's Deceptive And Misleading Omissions**

14 141. WF deceptively and misleadingly conceals other material facts about the Falsely
15 Labeled Products, including:

- 16 a. the true nature of the Falsely Labeled Products' ingredients;
- 17 b. the identity of the Falsely Labeled Products' ingredients;
- 18 c. that the Falsely Labeled Products contain sensitizers, irritants, toxins,
19 carcinogens, pollutants, and/or otherwise hazardous substances;
- 20 d. the concentration of the sensitizers, irritants, toxins, carcinogens, pollutants,
21 and/or otherwise hazardous substances in the Falsely Labeled Products;
- 22 e. that the Falsely Labeled Products are not "hypoallergenic";
- 23 f. that the Falsely Labeled Products are not what a reasonable consumer would
24 consider to be "hypoallergenic;"
- 25 g. that the Falsely Labeled Products contain chemicals that a reasonable
26 consumer would not expect in a product labeled as "hypoallergenic."

27 142. Plaintiff and the members of the Classes are not at fault for failing to discover WF's

1 wrongs earlier and had no actual or presumptive knowledge of facts sufficient to put them on
2 inquiry notice.

3 143. WF has concealed the identity of several ingredients. Discovery is therefore
4 necessary to determine their identity. These ingredients may also be sensitizers, irritants, or
5 otherwise toxic.

6 144. For example, WF adds “*fragrance*” or “*parfum*” to its products but does not
7 identify what chemical is used. Many ingredients used as fragrances are known skin sensitizers.
8 Many are also extremely toxic to a person’s skin, their overall health, and/or to the environment.

9 145. Furthermore, WF has not disclosed the concentration of each ingredient in its
10 products. Further investigation and discovery is needed so that Plaintiff can ascertain whether
11 entire products are also toxic.

12 146. WF has also concealed from consumers the nature of its products’ ingredients
13 despite consumers’ requests. The possible carcinogenic, toxic, and environmental effects of its
14 ingredients are still concealed from consumers today.

15 147. These facts are not ascertainable and are still not known to Plaintiff, the Class
16 members, and reasonable consumers. WF’s concealment tolls the applicable statute of limitations.

17 148. To this day, WF continues to conceal and suppress the existence, true identity,
18 nature, and concentration of the sensitizers, irritants, toxins, carcinogens, pollutants, and/or
19 otherwise hazardous substances in the Falsely Labeled Products.

20 149. Similarly, to this day, WF continues to conceal and suppress the fact that the Falsely
21 Labeled Products are not “hypoallergenic” as promised.

22 150. WF represents elsewhere on the product label and on its website that the products
23 are “non-toxic,” “safe,” having “only the gentlest ingredients,” and/or causing “no tears,” etc.
24 Exhibit 1. This further obscures the fact that WF’s products are not hypoallergenic.

25 151. For example, in its “Official Whole Foods Market Blog,” WF encourages
26 consumers seeking to avoid allergens in cleaning products to purchase Whole Foods Market brand
27 products, as they lack the ingredients WF identifies in its in-house list of banned “unacceptable

1 ingredients” for body care, premium body care, and household cleaners. *See, e.g.*, Exhibit 14
2 (“What You Won’t Find in our Cleaning Products”).

3 152. WF fails to disclose, however, that many ingredients in its products are known skin
4 allergens, even though they are not banned by WF’s list of “unacceptable ingredients.”

5 **G. WF Knew Its Representations Were False**

6 153. WF holds itself out to the public as trusted experts in the area of hypoallergenic,
7 safe, mild, and gentle personal care products.

8 154. WF knew what representations it made regarding the Falsely Labeled Products, as
9 all representations appear on the products’ packages.

10 155. WF also knew what ingredients were added to each product, as (presumably) all
11 product ingredients listed on the product packages and are further disseminated on their websites.

12 156. WF is governed by and thus is presumed to know the federal regulations and state
13 laws that control the labeling of the Falsely Labeled Products, and thus is aware that many of the
14 ingredients have been federally declared to be chemical compounds that require inventory
15 reporting under the Toxic Substance Control Act, are hazardous or toxic compounds that require
16 special disclosures on safety data sheets, or are carcinogens or reproductive toxins that require
17 product label warnings under state law.

18 157. WF thus knew all the facts demonstrating that its Falsely Labeled Products contain
19 sensitizers, irritants, and otherwise toxic ingredients, and that these products were therefore falsely
20 labeled.

21 **H. WF Intended Consumers To Rely**

22 158. As WF knows, consumers prefer hypoallergenic products. As WF knows,
23 consumers will pay a premium for hypoallergenic products or would not purchase these products
24 at all unless they were hypoallergenic, as advertised.

25 159. WF encourages consumers’ preference for hypoallergenic products – specifically
26 for WF’s products – explaining to consumers that “we believe the quality of the items and
27 ingredients you put on your body is as important as the foods and nutritional supplements you put

1 in your body.” Exhibit 12 (“Body Care Quality Standards”).

2 160. WF’s misleading affirmative statements (*e.g.*, that the products were mild, gentle,
3 safe, caused “no more tears,” or were environmentally safe) further obscured what WF failed to
4 disclose. Thus, reliance upon WF’s misleading and deceptive representations and omissions may
5 be presumed.

6 161. WF made the false, deceptive, and misleading representations and omissions,
7 intending Plaintiff and Class members to rely upon these representations and omissions in
8 purchasing and using one or more Falsely Labeled Products.

9 162. In making the false, misleading, and deceptive representations and omissions at
10 issue, WF knew and intended that consumers would purchase the WF products when consumers
11 would otherwise purchase a competing product or employ an alternate regimen (such as using an
12 oil for moisturizing).

13 163. In making the false, misleading, and deceptive representations and omissions at
14 issue, WF also knew and intended that consumers would pay a premium for hypoallergenic
15 products, furthering WF’s private interest of increasing sales of its products and decreasing the
16 sales of products marketed by its competitors.

17 **I. Consumers Reasonably Relied**

18 164. Consumers frequently rely on ingredient representations and information in making
19 purchase decisions, especially in purchasing personal care products.

20 165. When Plaintiff and the Class members purchased the Falsely Labeled Products,
21 Plaintiff and the Class members saw the false, misleading, and deceptive representations detailed
22 above, and did not receive disclosure of the facts concealed, as detailed above.

23 166. These misrepresentations were uniform and were communicated to Plaintiff and
24 every other member of the Class at every point of purchase and consumption.

25 167. Plaintiff and the Class members were among the intended recipients of WF’s
26 deceptive representations and omissions.

27 168. Plaintiff and the Class members reasonably relied to their detriment on WF’s

1 misleading representations and omissions.

2 169. WF's false, misleading, and deceptive misrepresentations and omissions deceived
3 and misled, and are likely to continue to deceive and mislead, Plaintiff, the Class members,
4 reasonable consumers, and the general public.

5 170. WF's misleading affirmative statements further obscured what it failed to disclose.
6 Thus, reliance upon WF's misleading and deceptive representations and omissions may be
7 presumed.

8 171. WF made the deceptive representations and omissions with the intent to induce
9 Plaintiff and the Class members to purchase the Falsely Labeled Products. Plaintiff and the Class
10 members' reliance upon such representations and omissions may be presumed.

11 172. WF's deceptive representations and omissions are material in that a reasonable
12 person would attach importance to such information and would be induced to act upon such
13 information in making purchase decisions. Thus, Plaintiff and the Class members' reliance upon
14 such representations and omissions may be presumed as a matter of law. The materiality of those
15 representations and omissions also establishes causation between WF's conduct and the injuries
16 sustained by Plaintiff and the Class members.

17 **J. WF's Wrongful Conduct Caused Plaintiff's Injury**

18 173. As an immediate, direct, and proximate result of WF's false, misleading, and
19 deceptive representations and omissions, WF injured Plaintiff and the Class members in that they:

- 20 a. paid a sum of money for a product that was not as represented;
- 21 b. paid a premium price for a product that was not as represented;
- 22 c. were deprived the benefit of the bargain because the Falsely Labeled
Products they purchased were different from what WF warranted;
- 24 d. were deprived the benefit of the bargain because the Falsely Labeled
Products they purchased had less value than what was represented;
- 26 e. did not receive a product that measured up to their expectations as created
by WF;

f. used (or caused their children to use) a substance that Plaintiff and the members of the Class did not expect or consent to;

g. used (or caused their children to use) a product that was not hypoallergenic;

h. without their knowing consent, used (or caused their children to use) a substance that is generally harmful to their health or their children's health;

i. without their knowing consent, used (or caused their children to use) a substance that is a skin sensitizer, irritant, or a known or suspected toxin, carcinogen, mutagen, teratogen, environmental pollutant, or otherwise is harmful to the environment and/or their health.

174. Had WF not made the false, misleading, and deceptive representations and omissions, Plaintiff and the Class members would not have been injured as listed above. Accordingly, Plaintiff and the Class members have suffered injury in fact as a result of WF's wrongful conduct.

175. Plaintiff and the Class members all paid money for the Falsely Labeled Products but did not obtain the full value of the advertised products due to WF's misrepresentations and omissions. Plaintiff and the Class members purchased, purchased more of, or paid more for, the Falsely Labeled Products than they would have had they known the truth about the Falsely Labeled Products. Accordingly, Plaintiff and the Class members have suffered injury in fact and lost money or property as a result of WF's wrongful conduct.

K. WF Benefitted From Its Misleading And Deceptive Representations And Omissions

176. As the intended, direct, and proximate result of WF's false, misleading, and deceptive representations and omissions, WF has been unjustly enriched through more sales of Falsely Labeled Products and higher profits at the expense of Plaintiff and the Class members. As a direct and proximate result of its deception, WF also unfairly obtained other benefits, including the higher value associated with a "hypoallergenic" brand and the resulting higher stock value, redirecting sales to it and away from its competitors, and increased sales of its other products.

IV. CLASS ALLEGATIONS

177. Plaintiff Kellman brings this action pursuant to Rule 23 of the Federal Rules of

1 Civil Procedure on behalf of themselves and all other similarly situated United States residents
2 who purchased the Falsely Labeled Products (as defined herein) (the “Nationwide Class”).

3 178. Plaintiff Kellman also brings this action on behalf of herself and all other similarly
4 situated California residents who purchased the Falsely Labeled Products (as defined herein) (the
5 “California Class”).

6 179. Excluded from the Class are the judge(s) assigned to this case and their family
7 members; officers and directors of WF; members of the immediate families of the officers and
8 directors of WF; WF’s legal representatives, heirs, successors, or assigns; and, any entity in which
9 they have or have had a controlling interest.

10 180. Plaintiff brings each Class pursuant to Federal Rules of Civil Procedure 23(a),
11 23(b)(1), 23(b)(2), and 23(b)(3).

12 181. At this time, Plaintiff does not know the exact number of the Class members; given
13 the nature of the claims and the number of sales that WF has made of the Products, Plaintiff
14 believes that members of each Class are so numerous that joinder of all members is impracticable.

15 182. There is a well-defined community of interest in the questions of law and fact
16 involved in this case. Questions of law and fact common to the members of the Class that
17 predominate over questions that may affect individual Class members include:

18 a. whether WF misrepresented and/or failed to disclose material facts
19 concerning the Falsely Labeled Products;

20 b. whether WF’s conduct was unfair and/or deceptive; and

21 c. whether WF breached an express warranty created through the labeling and
22 marketing of its Falsely Labeled Products.

23 183. Plaintiff’s claims are typical of those of the Class because Plaintiff, like all
24 members of the Class, purchased one or more of WF’s Falsely Labeled Products at a premium
25 price, relying on WF’s false and misleading representations, and Plaintiff sustained damages from
26 WF’s wrongful conduct.

27 184. Plaintiff will fairly and adequately protect the interests of the Class because

1 Plaintiff is similarly situated with, and have suffered similar injuries as, the members of the Class
 2 they seek to represent. Plaintiff feels that she has been deceived, wishes to obtain redress of the
 3 wrong, and wants WF to be stopped from perpetrating similar wrongs on others. Plaintiff is an
 4 adequate representative of the Class because her interests do not conflict with the interests of the
 5 Class members she seeks to represent, and she has retained counsel competent and experienced in
 6 conducting complex class action litigation, who were the first to publicly uncover the true scope
 7 and extent of WF's wrongs. Plaintiff has no interests adverse to those of the Class members and
 8 will vigorously prosecute this litigation.

9 185. A class action is superior to other available methods for the fair and efficient
 10 adjudication of this controversy. Specifically, no Class member has a substantial interest in
 11 individually controlling the prosecution of a separate action. The damages suffered by each
 12 individual Class member likely will be relatively small, especially given the burden and expense
 13 of individual prosecution of the complex litigation necessitated by WF's conduct. Thus, it would
 14 be virtually impossible for the Class members individually to redress effectively the wrongs done
 15 to them.

16 186. The prerequisites to maintaining a class action for injunctive or equitable relief are
 17 met as WF has acted or refused to act on grounds generally applicable to the Class, thereby making
 18 appropriate final injunctive or equitable relief with respect to the Class as a whole.

19 187. Upon information and belief, there are no pending lawsuits concerning the products
 20 at issue in this case. Concentration of the litigation concerning this matter in this Court is desirable,
 21 and the difficulties likely to be encountered in the management of a class action are not great. The
 22 resolution of the claims of all Class members in a single forum, and in a single proceeding, would
 23 be a fair and efficient means of resolving the issues raised in this litigation.

24 188. The prosecution of separate actions by Class members would create a risk of
 25 establishing inconsistent rulings and/or incompatible standards of conduct for WF.

26 189. WF's conduct is generally applicable to the Class as a whole and Plaintiff seeks,
 27 *inter alia*, equitable remedies with respect to the Class as a whole. As such, WF's systematic

1 policies and practices make declaratory relief with respect to the Class as a whole appropriate.

2 190. The Class is specifically identifiable to facilitate provision of adequate notice and
3 there will be no significant problems managing this case as a class action. Notice to the Class can
4 be made through various means, such as in-store leaflets, website notices, Facebook notices,
5 notices on the labels of the packages, and/or direct notice to those consumers for which WF knows
6 the e-mail or physical mailing address.

V. CAUSES OF ACTION

8 191. The allegations in each Cause of Action are repeated and realleged in every other
9 Cause of Action as if set forth in full therein.

COUNT 1

Breach of Express Warranty

***On Behalf of the Nationwide Class and, in the alternative,
the California Class***

192. WF provided Plaintiff and other members of the Class with written express
14 warranties including, but not limited to, warranties that its Falsely Labeled Products were
15 “hypoallergenic.”

193. These affirmations of fact or promises by WF relate to the goods and became part
17 of the basis of the bargain.

18 194. Plaintiff and members of each Class purchased the Falsely Labeled Products,
19 believing them to conform to the express warranties.

195. WF breached these warranties. This breach resulted in damages to Plaintiff and
other members of the Class, who bought Falsely Labeled Products but did not receive the goods
as warranted.

196. As a proximate result of the breach of warranties by WF, Plaintiff and the other
members of the Class did not receive goods as warranted. Plaintiff and the members of the Class
therefore have been injured and have suffered damages in an amount to be proven at trial. Among
other things, Plaintiff and members of the Class did not receive the benefit of the bargain and have

1 suffered other injuries as detailed above. Moreover, had Plaintiff and the Class members known
2 the true facts, they would not have purchased the products, would have purchased fewer products,
3 or would not have been willing to pay the premium price WF charged for the products.

4 WHEREFORE, Plaintiff prays for relief as set forth below.

5 **COUNT 2**

6 **Unjust Enrichment**

7 ***On Behalf of the Nationwide Class and, in the alternative,
the California Class***

8 197. As a result of WF's deceptive, fraudulent, and misleading labeling, advertising,
9 marketing, and sales of the Falsely Labeled Products, WF was enriched at the expense of Plaintiff
10 and the other members of the Class through the payment of the purchase price for WF's Falsely
11 Labeled Products.

12 198. Under the circumstances, it would be against equity and good conscience to permit
13 WF to retain the ill-gotten benefits that it received from Plaintiff and the other members of the
14 Class, in light of the fact that the Falsely Labeled Products purchased by Plaintiff and the other
15 members of the Class were not what WF purported them to be. Thus, it would be unjust or
16 inequitable for WF to retain the benefit without restitution to Plaintiff and the other members of
17 the Class for the monies paid to WF for such Falsely Labeled Products.

18 WHEREFORE, Plaintiff prays for relief as set forth below.

19 **COUNT 3**

20 **Unfair and Deceptive Acts and Practices**

21 ***On Behalf of the Nationwide Class and, in the alternative, the California Class***

22 199. This cause of action is brought pursuant to California's Consumers Legal Remedies
23 Act, Cal. Civ. Code §§ 1750-1785 ("CLRA") and similar statutes.

24 200. Plaintiff and the other members of the Class are "consumers," as the term is defined
25 by California Civil Code § 1761(d) and similar statutes, because they bought the Falsely Labeled
26 Products for personal, family, or household purposes. WF is a "person" under Cal. Civ. Code
27

1 § 1761(c) and similar statutes.

2 201. The Falsely Labeled Products are “goods” under Cal. Civ. Code § 1761(a) and
3 similar statutes. Plaintiff, the other members of the Class, and WF have engaged in “transactions,”
4 as that term is defined by California Civil Code § 1761(e) and similar statutes. For the California
5 Class, these transactions all occurred in the State of California.

6 202. The conduct alleged in this Complaint constitutes unfair methods of competition
7 and unfair and deceptive acts and practices for the purposes of the CLRA and similar statutes, and
8 the conduct was undertaken by WF in transactions intended to result in, and which did result in,
9 the sale of goods to consumers.

10 203. WF’s false and fraudulent representations and omissions have violated and
11 continue to violate the CLRA and similar statutes because they extend to transactions that are
12 intended to result, or have resulted, in the sale of goods to consumers, including the Plaintiff and
13 the Class members.

14 204. WF’s conduct violates Cal. Civ. Code § 1770(a)(5) and similar statutes, which
15 prohibits “[r]epresenting that goods . . . have . . . characteristics [or] ingredients . . . which they do
16 not have,” and Cal. Civ. Code § 1770(a)(7) and similar statutes, which prohibits: “[r]epresenting
17 that goods . . . are of a particular standard, quality, or grade . . . if they are of another,” causing
18 injury to Plaintiff and the Class.

19 205. As a result of engaging in such conduct, WF has violated California Civil Code
20 § 1770(a)(5), (a)(7), and (a)(9) and similar statutes.

21 206. Plaintiff served WF with notice of its CLRA violations by certified mail, return
22 receipt requested.

23 207. Plaintiff and the Class members seek preliminary injunctive relief, and permanent
24 injunctive relief against WF’s unfair and deceptive acts and conduct.

25 208. Pursuant to California Civil Code § 1780(a)(2) and (a)(5) and similar statutes,
26 Plaintiff seeks an order of this Court that includes, but is not limited to, an order enjoining WF
27 from continuing to engage in unlawful, unfair, or fraudulent business practices or any other act

1 prohibited by law.

2 209. Plaintiff and the other Class members may be irreparably harmed and/or denied an
3 effective and complete remedy if such an order is not granted.

4 210. The unfair and deceptive acts and practices of WF, as described above, present a
5 serious threat to Plaintiff and the other members of the Class.

6 WHEREFORE, Plaintiff prays for relief as set forth below.

7 **COUNT 4**

8 **Violations of California's False Advertising Law and Similar Statutes**

9 ***On Behalf of the Nationwide Class and, in the alternative, the California Class***

10 211. This cause of action is brought pursuant to California's False Advertising Law
11 ("FAL"), Cal. Bus. & Prof. Code § 17500 *et seq.* and similar statutes.

12 212. Such acts of WF, as described above, and each of them constitute unlawful,
13 deceptive, and fraudulent business acts and practices.

14 213. At all material times, WF engaged in a scheme of offering the Falsely Labeled
15 Products for sale to Plaintiff and the other members of the Class by way of distributing within the
16 State of California (or the residence) to the public, *inter alia*, commercial marketing and
17 advertising, the World Wide Web (Internet), Falsely Labeled Product packaging and labeling, and
18 other promotional materials and offered for sale the Falsely Labeled Products on a nationwide
19 basis, including in California.

20 214. The misrepresentations and non-disclosures by WF of the material facts detailed
21 above constitute false and misleading advertising, and therefore constitute a violation of Cal. Bus.
22 & Prof. Code § 17500, *et seq.* and similar statutes.

23 215. Said advertisements and inducements were made within the state of residence and
24 come within the definition of advertising contained in the FAL in that such promotional materials
25 were intended as inducements to purchase WF's Falsely Labeled Products and are statements
26 disseminated by WF to Plaintiff and the other Class members. WF knew, or in the exercise of
27 reasonable care should have known, that these representations were misleading and deceptive.

1 216. Consumers, including Plaintiff and the other Class members, necessarily and
2 reasonably relied on these materials concerning WF's Falsely Labeled Products. Consumers,
3 including Plaintiff and the Class members, were among the intended targets of such
4 representations.

5 217. The above acts of WF did and were likely to deceive reasonable consumers,
6 including Plaintiff and the other members of the Class, by obfuscating the nature, quality, and/or
7 ingredients of the Falsely Labeled Products, in violation of the “misleading” prong of the FAL and
8 similar statutes.

9 218. The business practices alleged above are unlawful under the CLRA and similar
10 statutes, which forbids misleading and deceptive advertising.

11 219. Plaintiff and the other members of the Class have suffered injury in fact and have
12 lost money or property as a result of WF's violations of the FAL and similar statutes.

13 220. As a result, WF has been unjustly enriched at the expense of Plaintiff and the other
14 members of the Class. Plaintiff and the Class, pursuant to California Business and Professions
15 Code § 17535 and similar statutes, are entitled to an order of this Court enjoining such future
16 conduct on the part of WF, and such other orders and judgments which may be necessary to
17 disgorge WF's ill-gotten gains and restore to any person in interest any money paid for its Falsely
18 Labeled Products as a result of the wrongful conduct of WF.

19 WHEREFORE, Plaintiff prays for relief as set forth below.

COUNT 5

Violation of California's Unfair Competition Law and Similar Statutes

On Behalf of the Nationwide Class and, in the alternative, the California Class

23 221. This cause of action is brought pursuant to California's Unfair Competition Law
24 ("UCL"), Cal. Bus. & Prof. Code § 17200 *et seq.* and similar statutes.

25 222. By committing the acts and practices alleged herein, WF has engaged in deceptive,
26 unfair, and unlawful business practices in violation of the UCL and similar statutes.

223. Plaintiff has standing to pursue this claim as she has suffered injury in fact and has

1 lost money or property as a result of WF's actions as set forth above. Class members also have
2 suffered injury in fact and have lost money or property as a result of WF's actions as set forth
3 above.

4 224. The violation of any law constitutes an "unlawful" business practice under Cal.
5 Bus. & Prof. Code § 17200 and similar statutes.

6 225. Each of WF's false representations alleged herein violates U.S.C. § 331; Cal. Civ.
7 Code § 1709; Cal. Civ. Code § 1750 *et seq.*; and Cal. Bus. & Prof. Code § 17500 *et seq.*, and
8 similar statutes.

9 226. WF has violated the UCL's proscription against engaging in unlawful conduct as a
10 result of its violations of (i) the CLRA and similar statutes, as alleged above, and (ii) the FAL and
11 similar statutes, as alleged above.

12 227. In addition, WF has violated the UCL's proscription against engaging in unlawful
13 conduct as a result of its violations of the Sherman Law, Cal. Health & Safety Code § 109875 *et*
14 *seq.*, and similar statutes, which forbid (1) misbranding of any cosmetic, *id.* at §§ 110398 and
15 111445, and (2) manufacturing, selling, delivering, holding, or offering for sale any cosmetic that
16 is misbranded or delivering or proffering such for delivery. Cal. Health & Safety Code §§ 110390,
17 110395, 110398, 110400, 110550, 110585, 110620, 110625, 110660, 110770, 110705, 110740,
18 110760, 110765, 110770, 111445, and 111450.

19 228. The Sherman Law defines a "person" as "any individual, firm, partnership, trust,
20 corporation, limited liability company, company, estate, public or private institution, association,
21 organization, group, city, county, city and county, political subdivision of this state, other
22 governmental agency within the state, and any representative, agent, or agency of any of the
23 foregoing." Cal. Health & Safety Code § 109995. WF is a "person" within the meaning of the
24 Sherman Law.

25 229. As more fully described herein, WF's misleading marketing, advertising,
26 packaging, and labeling of the Falsely Labeled Products is likely to deceive a reasonable consumer.
27 Indeed, Plaintiff and the other Class members were unquestionably deceived regarding the

1 characteristics of WF's Falsely Labeled Products, as WF's marketing, advertising, packaging, and
 2 labeling of the Falsely Labeled Products misrepresents and/or omits the true nature, quality, and/or
 3 ingredients of the Falsely Labeled Products.

4 230. There is no benefit to consumers or competition from deceptively marketing and
 5 labeling products. Indeed, the harm to consumers and competition is substantial. Plaintiff and the
 6 other members of the Class who purchased the Falsely Labeled Products suffered a substantial
 7 injury as alleged herein.

8 231. Plaintiff and the other members of the Class who purchased the Falsely Labeled
 9 Products had no way of reasonably knowing that the Falsely Labeled Products they purchased
 10 were not as marketed, advertised, packaged, and labeled. Thus, they could not have reasonably
 11 avoided the injury each of them suffered.

12 232. WF's acts and omissions alleged above constitute unfair business practices under
 13 Cal. Bus. & Prof. Code § 17200 and similar statutes because the gravity of the consequences of
 14 WF's conduct as described above outweighs any justification, motive, or reason therefor,
 15 particularly considering the available legal alternatives which exist in the marketplace, and such
 16 conduct is immoral, unethical, unscrupulous, offends established public policy, or is substantially
 17 injurious to Plaintiff and the other members of the Class. WF's false and misleading
 18 representations and omissions also violate legislatively declared policy as they have violated
 19 numerous state and federal laws. Moreover, the gravity of the harm to Plaintiff and Class members
 20 resulting from WF's conduct outweighs WF's legitimate reasons, justifications and/or motives for
 21 engaging in such deceptive acts and practices

22 233. Each false and misleading representation and omission constitutes fraudulent
 23 business practices under Cal. Bus. & Prof. Code § 17200 and similar statutes because the
 24 representations and omissions were false. Even if these representations were true, WF's
 25 representations and deceptive concealment were nonetheless fraudulent under the statute because
 26 they were misleading and were likely to and did deceive the reasonable consumer, including
 27 Plaintiff and the Class members.

1 234. WF's violations continue to this day.

2 235. Pursuant to California Business and Professions Code § 17203 and similar statutes,
3 Plaintiff and the other members of the Class seek an order of this Court that includes, but is not
4 limited to, an order enjoining such future conduct on the part of WF and such other orders and
5 judgments which may be necessary to disgorge WF's ill-gotten gains and to restore to any person
6 in interest any money paid for WF's Falsely Labeled Products as a result of the wrongful conduct
7 of WF.

8 WHEREFORE, Plaintiff prays for relief as set forth below.

9 **PRAYER FOR RELIEF**

10 WHEREFORE, Plaintiff demands judgment on behalf of herself and the proposed Classes
11 providing such relief as follows:

12 A. Certification of the Classes proposed herein under Federal Rule of Civil
13 Procedure 23(a), (b)(1), (b)(2), and (b)(3); appointment of Plaintiff Kellman as representative of
14 the Nationwide Class and the California Class; and, appointment of the undersigned counsel as
15 counsel for the Classes;

16 B. A declaration that WF is financially responsible for notifying members of the
17 Classes of the pendency of this suit;

18 C. An order requiring an accounting for, and imposition of a constructive trust
19 upon, all monies received by WF as a result of the unfair, misleading, fraudulent, and unlawful
20 conduct alleged herein;

21 D. Restitution, disgorgement, refund, and/or other monetary damages, together
22 with costs and disbursements, including reasonable attorneys' fees pursuant to the applicable
23 statutes and prejudgment interest at the maximum rate allowable by law;

24 E. Injunctive relief on behalf of the Classes, enjoining WF's unlawful and
25 deceptive acts;

26 F. Statutory damages in the maximum amount provided by law;

27 G. Punitive damages in accordance with proof and in an amount consistent with

1 applicable precedent; and

2 H. Such further relief as this Court may deem just and proper.

3 **JURY TRIAL DEMANDED**

4 Plaintiff and the Class members hereby demand a trial by jury.

5
6 DATED: January 2, 2019

Stephanie R. Tatar – State Bar No. 237792
TATAR LAW FIRM, APC
3500 West Olive Avenue, Suite 300
Burbank, California 91505
Tel. (323) 744-1146
Fax. (888) 778-5695
Stephanie@thetatarlawfirm.com

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8
9
10 **THE GOLAN FIRM**

11 Yvette Golan (*pro hac vice*)
12 1712 N Street, NW, Suite 302
Washington, D.C. 20036
13 Tel: (866) 298-4150, ext. 101
Fax: (928) 441-8250

14
15 s/ James A. Francis

16 **FRANCIS & MAILMAN, P.C.**
17 James A. Francis (*pro hac vice*)
David A. Searles (*pro hac vice*)
18 1600 Market Street, Suite 2510
Philadelphia, PA 19103
Tel. (215) 735-8600
Fax. (215) 950-8000